This listing of claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims:

- (Currently Amended): <u>A process</u> Process for the reductive preparation of nicotinaldehydes a nicotinicaldehyde, said process comprising performing reduction on characterised in that the starting materials employed for the reduction are the corresponding nicotinic acid <u>morpholinamide</u> in the presence of a reducing agent morpholinamides to obtain said nicotinicaldehyde.
- (Currently Amended): <u>A process</u> Process according to Claim 1, characterised in that the starting materials employed are <u>wherein said</u> nicotinic acid <u>morpholinamide is</u> morpholinamides of the formula I

## wherein in which

- $R^{1}$ ,  $R^{1o}$   $\underline{are}$  each, independently of one another, <del>denotes</del> H, Hal, A, OA,  $CH_2R^2$  or Ar,  $R^2$   $\underline{ig}$  denotes OA or NA<sub>2</sub>,
- A is denotes unbranched or branched alkyl having 1-10 C atoms, in which one or two CH<sub>2</sub> groups are each optionally may be replaced by an O or S atom or atoms and/or-by a -CH=CH- group groups and/or also 1-7 H atoms are each optionally may be replaced by F, or A is cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclocytyl, 2.6.6-trimethylbicyclo[3.1.1]heptyl, p-menthane, menthol, pinane, bornane, camphor, or adamantly.
- Ar <u>is denotes</u> an unsaturated, partially or fully saturated, mono- or polycyclic homo- or heterocyclic system <u>wherein</u> with the hetero atoms <u>are each O, N, or S, and</u> which is unsubstituted or mono- or polysubstituted by Hal, A, OA, NA<sub>2</sub>, NO<sub>2</sub>, NASO<sub>2</sub>A, SO<sub>2</sub>NA, SO<sub>2</sub>A, and

## Hal is denotes F, Cl, Br or I.

- (Currently Amended): <u>A process</u> Process according to Claim I, characterised in that the starting material employed wherein said nicotinic acid morpholinamide is 5-(4fluorophenyl)nicotinic acid morpholinamide.
- (Currently Amended): <u>A process</u> Process according to Claim I, characterised in that the starting material employed wherein said nicotinic acid morpholinamide is 5-bromopyridine-3-carboxylic acid morpholinamide.
- (Currently Amended): <u>A process</u> Process according to Claim 1, eharacterised in that the <u>wherein said</u> reducing <u>agent is agents-employed are LiAlH(OEt)</u>, LiAlH<sub>2</sub>(OEt)<sub>2</sub> or LiAlH<sub>3</sub>(OEt).
  - (Cancelled):
  - (Cancelled):
- (Currently Amended): <u>A process</u> <u>Starting materials of the formula I</u> according to Claim I, <u>wherein said nicotinic acid morpholinamide is selected from a group consisting of</u>
  (a) 5-(4-fluorophenyl)nicotinic acid morpholinamide <u>or</u> [[,]] (b) 5-bromonicotinic acid morpholinamide.
- 9. (New): A process according to Claim 1, wherein the reducing agent for said reduction is  $LiAlH_{(4-n)}(OR)_n$ , where n is 1, 2 or 3, and R in each case is methyl, ethyl or tertbutyl.
- (New): A process according to Claim 2, wherein R<sup>V</sup>, R<sup>V</sup> are each, independently of one another, hydrogen, methoxy, ethoxy, propoxy, butoxy, fluorine, chlorine, bromine, iodine, phenyl, or o, m or p-substituted phenyl.

- 11. (New): A process according to Claim 2, wherein  $R^{\Gamma}$  is p-fluorophenyl or bromine and  $R^{\Gamma}$  is hydrogen.
- (New): A process according to Claim 2, wherein Hal is fluorine, chlorine or bromine.
- 13. (New): A process according to Claim 2, wherein A is unbranched or branched alkyl having 1-10 C atoms, in which one or two CH<sub>2</sub> groups are each optionally replaced by an O or S atom or by a -CH=CH- group and/or also 1-7 H atoms are each optionally replaced by F.
- 14. (New): A process according to Claim 13, wherein A is methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, 1-, 2- or 3-methylbutyl, 1,1-, 1,2- or 2,2-dimethylpropyl, 1-ethylpropyl, hexyl, 1-, 2-, 3- or 4-methylpentyl, 1,1-, 1,2-, 1,3-, 2,2-, 2,3- or 3,3-dimethylbutyl, 1- or 2-ethylbutyl, 1-ethyl-1-methylpropyl, 1-ethyl-2-methylpropyl, 1,1,2- or 1,2,2-trimethylpropyl, or trifluoromethyl.
- (New): A process according to Claim 13, wherein A is methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, trifluoromethyl, pentafluoroethyl, or 1,1,1-trifluoroethyl.
- (New): A process according to Claim 2, wherein Ar is unsubstituted or substituted phenyl, naphthyl or biphenyl.
- 17. (New): A process according to Claim 2, wherein Ar is phenyl, o-, m- or p-tolyl, o-, m- or p-cyanophenyl, o-, m- or p-methoxyphenyl, o-, m- or p-fluorophenyl, o-, m- or p-bromophenyl, o-, m- or p- chlorophenyl, 2,3-, 2,4-, 2,5-, 2,6-, 3,4- or 3,5-difluorophenyl, 2,3-, 2,4-, 2,5-, 2,6-, 3,4- or 3,5-dibromophenyl, 2,3-, 2,4-, 2,5-, 2,6-, 3,4- or 3,5-dibromophenyl, 2-fluoro-4-bromophenyl, or 2,5-difluoro-4-bromophenyl.
  - 18. (New): A process according to Claim 2, wherein

- R<sup>1</sup>, R<sup>1</sup> are each, independently of one another, hydrogen, methoxy, ethoxy, propoxy, butoxy, fluorine, chlorine, bromine, iodine, phenyl, or o, m or p-substituted phenyl;

  Hal is fluorine chlorine or bromine:
- A is methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, 1-, 2- or 3-methylbutyl, 1,1-, 1,2- or 2,2-dimethylpropyl, 1-ethylpropyl, hexyl, 1-, 2-, 3- or 4-methylpentyl, 1,1-, 1,2-, 1,3-, 2,2-, 2,3- or 3,3-dimethylbutyl, 1- or 2-ethylbutyl, 1-ethyl-1-methylpropyl, 1-ethyl-2-methylpropyl, 1,1,2- or 1,2,2-trimethylpropyl, trifluoromethyl, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cyclohetyl, cycloctyl, 2,6,6-trimethylbicyclo[3.1.1]heptyl, p-menthane, menthol, pinane, bornane, camphor, or adamantly; and
- Ar is phenyl, o-, m- or p-tolyl, o-, m- or p-cyanophenyl, o-, m- or p-methoxyphenyl, o-, m- or p-fluorophenyl, o-, m- or p-bromophenyl, o-, m- or p- chlorophenyl, 2,3-, 2,4-, 2,5-, 2,6-, 3,4- or 3,5-difluorophenyl, 2,3-, 2,4-, 2,5-, 2,6-, 3,4- or 3,5-difluorophenyl, 2,3-, 2,4-, 2,5-, 2,6-, 3,4- or 3,5-difluorophenyl, 2-fluoro-4-bromophenyl, or 2,5-difluoro-4-bromophenyl.
- 19. (New): A process according to Claim 2, wherein the reduction is carried out in an inert solvent, and said solvent is selected from hexane, petroleum ether, benzene, toluene or xylene; ethers, such as diethyl ether, diisopropyl ether, tetrahydrofuran, dioxane, ethylene glycol dimethyl ether, and mixtures of thereof.
- (New): A process according to Claim 19, wherein the amount of solvent is 10 g to 100 g of solvent per g of nicotinic acid morpholinamide.
- (New): A process according to Claim 2, wherein the reduction is performed at a temperature between -10° and 100°.
- (New): A process according to Claim 19, wherein the reduction is performed at a temperature between -10° and 100°.